

**K850057 DU PONT TOBRAMYCIN ASSAY CALIBRATOR**Jan 18, 1985  
10 days to decisionK850057 · Product code: **DLJ** · Toxicology  
Source: <https://www.510kdatabase.net/k850057/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Specific (DLJ)
Date received	Jan 8, 1985
Decision date	Jan 18, 1985
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>E.I. Dupont DE Nemours &amp; Co., Inc.</b>
Location	Mchenry, IL, US
Contact	ROGER C BRIDEN
510(k) history	253 submissions · 252 cleared · 1976-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k850057/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 14, 2026